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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/037,516	01/04/2002	Ashkan Imanzahrai	31505.0001	6624

7590

11/20/2002

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EXAMINER

DELACROIX MUIRHEI, CYBILLE

ART UNIT PAPER NUMBER

1614

DATE MAILED: 11/20/2002

8

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary

Application No.

10/037,516

Applicant(s)

IMANZAHRAL

Examiner

Cybill Delacroix-Muirheid

Art Unit

1614

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 26 August 2002.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 16, 18, 20, 22 and 26 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☒ Claim(s) 18, 20 and 22 is/are allowed.
- 6) ☒ Claim(s) 16 and 26 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
- Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- 11) ☐ The proposed drawing correction filed on _____ is: a) ☐ approved b) ☐ disapproved by the Examiner.
- If approved, corrected drawings are required in reply to this Office action.
- 12) ☐ The oath or declaration is objected to by the Examiner.

Priority under 35 U.S.C. §§ 119 and 120

- 13) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
- ☐ Certified copies of the priority documents have been received.
 - ☐ Certified copies of the priority documents have been received in Application No. _____.
 - ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- * See the attached detailed Office action for a list of the certified copies not received.
- 14) ☒ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).
- a) ☐ The translation of the foreign language provisional application has been received.
- 15) ☒ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

Attachment(s)

- 1) ☐ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☒ Information Disclosure Statement(s) (PTO-1449) Paper No(s) Z.
- 4) ☐ Interview Summary (PTO-413) Paper No(s). _____.
- 5) ☐ Notice of Informal Patent Application (PTO-152)
- 6) ☐ Other: _____.

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DETAILED ACTION

The following is responsive to Applicant's amendment received Aug. 28, 2002.

Claims 2, 4, 6, 8, 10, 12, 14, 24, 28, 31, 34, 36, 38, 41 and 42 are cancelled.

No new claims are added.

Claims 16, 18, 20, 22, 26 are currently pending.

The indicated allowability of claims 16 and 26 is withdrawn in view of the following new ground of rejection.

The previous claims objections set forth in paragraphs 1-2 of the office action mailed March 13, 2002 are withdrawn in view of Applicant's amendment and the remarks contained therein.

The previous claim rejection under 35 USC 112, paragraph 2, set forth in paragraph 3 of the office action mailed March 13, 2002, is withdrawn in view of Applicant's amendment and the remarks contained therein.

The previous claims rejection under 35 USC 103(a) set forth in paragraphs 4-6 of the office action mailed March 13, 2002 is withdrawn in view of Applicant's amendment and the remarks contained therein.

Information Disclosure Statement

Applicant's Information disclosure Statement received Oct. 17, 2002 has been considered.

Please refer to Applicant's copy of the 1449 submitted herewith.

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Claim Objections

1. Claim 26 is objected to because of the following informalities: in claim 26, line 2, the term “migrain” should read --migraine--. Appropriate correction is required.

Claim Rejections - 35 USC § 112

2. Claim 26 is rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

In claim 26, lines 1-3, the phrase “an oral composition to treat migraine pain and other associated symptoms comprising a medicament component consisting of ...” renders the claim indefinite because it is not clear as to what components are excluded from and/or included in the claimed composition. The metes and bounds of the patent protection desired is unclear. For purposes of this office action, the claim will be interpreted as an “open” claim.

Claim Rejections - 35 USC § 103

3. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

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4. The factual inquiries set forth in *Graham v. John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

1. Determining the scope and contents of the prior art.
2. Ascertaining the differences between the prior art and the claims at issue.
3. Resolving the level of ordinary skill in the pertinent art.
4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

5. Claims 16 and 26 are rejected under 35 U.S.C. 103(a) as being unpatentable over Armellino et al., 5,972,916 (already of record) in view of Munari et al. (as explained by Applicant's specification page 12, lines 9-23).

Armellino et al. teach a method and composition for treating migraine headaches and associated symptoms, i.e. nausea, phonophobia, photophobia, pain and functional disability, (see Applicant's claim 16, which recites "or functional disability" (in the alternative)) the method comprising administering an effective amount of a composition comprising acetaminophen, caffeine, aspirin and a pharmaceutically acceptable carrier such as a lubricants or disintegrants. The composition may be administered orally in tablet form or may administered in liquid form. The dosing interval is daily every four to six hours. Please see the abstract; col. 2-col. 3; col. 6, lines 3-56; col. 7, lines 30-33.

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Armellino et al. do not disclose a method or composition for treating migraines and associated symptoms, i.e. functional disability, where the method administers a composition additionally containing pseudoephedrine; however, the Examiner refers to the Munari et al. reference which discloses a study where pseudoephedrine was administered to migraine patients. The data demonstrated that pseudoephedrine was effective in treating “cardiovascular abnormalities”, namely lower blood pressure and postural hypotension (see page 375 and 377; also, see Applicant’s specification, page 12, lines 8-22.

It would have been obvious to one of ordinary skill in the art at the time the invention was made to modify the method and composition of Armellino et al. by combining pseudoephedrine with the acetaminophen-containing composition of Armellino et al. because one of ordinary skill in the art would reasonably expect the additive effect of the acetaminophen-containing compositions and the pseudoephedrine to be effective in treating migraine pain and functional disability associated therewith. Moreover, one of ordinary skill in the art would reasonably expect pseudoephedrine to treat any cardiovascular abnormalities, i.e. a “functional disability”, that the patients of Armellino may experience. Therefore, such a modification would have been motivated by the reasoned expectation of successfully and comprehensively treating a migraine and cardiovascular abnormalities associated therewith.

With respect to the claimed dosage amounts of pseudoephedrine, since Munari et al. establish that pseudoephedrine’s efficacy is dependent upon dosage amounts, it would have been obvious to one of ordinary skill in the art to further modify the dosage amounts in the prior art such that

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pseudoephedrine is administered in an amount which is effective to optimize its effect in the treatment of cardiovascular abnormalities associated with migraines. Finally, with respect to administering the acetaminophen and pseudoephedrine during the prodrome phase of a migraine attack, it would have been obvious to one of ordinary skill in the art to modify the timing of administration during a migraine attack such that optimum treatment of the migraine and associated symptoms is achieved.

In addressing Armellino's use of caffeine and aspirin in the methods and composition, Applicant is reminded that the instant claims, recite "comprising" language which opens the claims and does not exclude other ingredients taught by the prior art but not claimed by Applicant. The transitional term "comprising", which is synonymous with "including," "containing," or "characterized by," is inclusive or open-ended and does not exclude additional, unrecited elements or method steps. Moleculon Research Corp. v. CBS, Inc., 793 F.2d 1261, 229 USPQ 805 (Fed. Cir. 1986); In re Baxter, 656 F.2d 679, 686, 210 USPQ 795, 803 (CCPA 1981); Ex parte Davis, 80 USPQ 448, 450 (Bd. App.1948)("comprising" leaves "the claim open for the inclusion of unspecified ingredients even in major amounts"). Please see MPEP 2111.03.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Cybille Delacroix-Muirheid whose telephone number is (703) 306-3227. The examiner can normally be reached on Tue-Fri from 8:30 to 6:00. The examiner can also be reached on alternate Mondays.

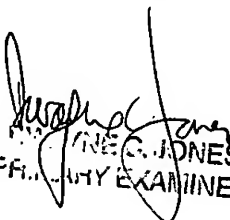
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If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Marianne Seidel, can be reached on (703) 308-4725. The fax phone number for this Group is (703) 308-4242.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the Group receptionist whose telephone number is (703) 308-1235.

CDM

Nov. 17, 2002


GREGORY JONES
PATENT EXAMINER